**UKHSA publications gateway number: GOV-18489**

## Shingles (Shingrix®) Herpes Zoster Vaccine Patient Group Direction (PGD)

This PGD is for the administration of Shingrix® herpes zoster vaccine (recombinant, adjuvanted), for the prevention of herpes zoster (‘zoster’ or shingles) and herpes zoster-related post-herpetic neuralgia (PHN), to individuals who are eligible for the national shingles immunisation programme.

This PGD is for the administration of Shingrix® by registered healthcare practitioners identified in [section 3](#_Characteristics_of_staff), subject to any limitations to authorisation detailed in [section 2](#section2).

Reference no: Shingles (Shingrix®) PGD

Version no:v3.0

Valid from: 1 September 2025

Review date: 29 February 2028

Expiry date: 31 August 2028

**The UK Health Security Agency (UKHSA) has developed this PGD to facilitate the delivery of publicly-funded immunisation in England in line with national recommendations.**

Those using this PGD must ensure that it is organisationally authorised and signed in Section 2 by an appropriate authorising person, relating to the class of person by whom the product is to be supplied, in accordance with Human Medicines Regulations 2012 (HMR2012)[[1]](#footnote-2). **The PGD is not legal or valid without signed authorisation in accordance with** [**HMR2012 Schedule 16 Part 2**](http://www.legislation.gov.uk/uksi/2012/1916/schedule/16/part/2/made)**.**

Authorising organisations must not alter, amend or add to the clinical content of this document (sections 4, 5 and 6); such action will invalidate the clinical sign-off with which it is provided. In addition, authorising organisations must not alter Section 3 (Characteristics of staff). **Sections 2 and 7 can be amended within the designated editable fields provided, but only for the purposes for which these sections are provided, namely the responsibilities and governance arrangements of the NHS organisation using the PGD. The fields in section 2 and 7 cannot be used to alter, amend or add to the clinical content. Such action will invalidate the UKHSA clinical content authorisation which is provided in accordance with the regulations**.

Operation of this PGD is the responsibility of commissioners and service providers. The final authorised copy of this PGD should be kept by the authorising organisation completing Section 2 for 8 years after the PGD expires as the PGD relates to adults only. Provider organisations adopting authorised versions of this PGD should also retain copies for 8 years.

**Individual practitioners must be authorised by name, under the current version of this PGD before working according to it.**

Practitioners and organisations must check that they are using the current version of the PGD. Amendments may become necessary prior to the published expiry date. Current versions of UKHSA PGD templates for authorisation can be found from: [Immunisation patient group direction (PGD) templates](https://www.gov.uk/government/collections/immunisation-patient-group-direction-pgd)

Any concerns regarding the content of this PGD should be addressed to:

immunisation@ukhsa.gov.uk

Enquiries relating to the availability of organisationally authorised PGDs and subsequent versions of this PGD should be directed to: Insert local contact details such as SIT inbox

# **Change history**

|  |  |  |
| --- | --- | --- |
| **Version number** | **Change details** | **Date** |
| v1.0 | New Shingrix® Herpes Zoster Vaccine PGD.  | 22 August 2021 |
| v2.0 | Shingrix® Herpes Zoster Vaccine PGD amended to include: * addition of new eligibility cohorts to reflect policy change, effective as of 1 September 2023 (see [Appendix 1](#Appendix1) for summary).

clarification of co-administration of Shingrix® with adjuvanted influenza vaccine. | 14 July 2023  |
| v3.0 | Shingles (Shingrix®)Herpes Zoster Vaccine PGD amended to include: * the addition of new eligibility cohorts, [reflecting policy change](https://www.gov.uk/government/publications/shingles-herpes-zoster-vaccination-programme-jcvi-statement-november-2024/jcvi-statement-on-the-shingles-herpes-zoster-vaccination-programme#expansion-of-eligibility-immunosuppressed-people) effective as of 1 September 2025
* an emphasis that eligible individuals with severe immunosuppression includes those who are due to commence treatment imminently
* an emphasis on the minimum recommended intervals between Shingrix® doses in England and appropriate action to take when shorter intervals are required
* remove references to Zostavax®, which was no longer in use in the UK beyond 31 October 2024
* an exception to repeat shingles vaccination for individuals who become severely immunosuppressed subsequent to having been immunised with Zostavax® in the past
* registered healthcare professionals named in both the GP Additional Roles Reimbursement Scheme (ARRS) and [HMR2012](https://www.legislation.gov.uk/uksi/2012/1916/schedule/16/part/4/made), namely dieticians, occupational therapists, pharmacy technicians and podiatrists
* updated written information for individuals and carers
* updated Appendix A
* minor formatting and other revisions to bring this PGD template in line with other UKHSA PGDs
 | 22 July 2025 |

1. **PGD development**

This PGD has been developed by the following health professionals on behalf of UKHSA:

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| **Developed by:** | **Name** | **Signature** | **Date** |
| Pharmacist(Lead author) | Christina WilsonLead Pharmacist, Immunisation Programmes Division, UKHSA |  | 21 July 2025 |
| Doctor | Dr Gayatri AmirthalingamDeputy Director of Public Health Programmes and Consultant Epidemiologist, Immunisation and Vaccine Preventable Diseases Division, UKHSA  |  | 21 July 2025 |
| Registered Nurse(Chair of Expert Panel) | David GreenNurse Consultant, Immunisation Programmes Division, UKHSA |  | 21 July 2025 |

This PGD has been peer reviewed by the UKHSA Immunisations PGD Expert Panel in accordance with the UKHSA PGD and Protocol Policy. It has been ratified by the UKHSA Medicines Governance Committee

**Expert Panel**

|  |  |
| --- | --- |
| **Name** | **Designation** |
| Dr Nicholas Aigbogun | Consultant in Communicable Disease Control, Yorkshire and Humber Health Protection Team, UKHSA  |
| Jessica Baldasera | Health Protection Practitioner, North East Health Protection Team, Regions Directorate, UKHSA  |
| Helen Beynon  | Clinical Advisor, Immunisation Clinical Advice Response Service (CARS), NHSE London |
| Alison Campbell | Screening and Immunisation Coordinator, Clinical, NHSE Midlands |
| Naveen Dosanjh | Senior Clinical Advisor – Vaccinations, NHS England  |
| Rosie Furner | Advanced Specialist Pharmacist, Medicines Governance (Patient Group Directions and Medicines Mechanisms), NHS Specialist Pharmacy Service |
| Ed Gardner | Advanced Paramedic Practitioner / Emergency Care Practitioner, Medicines Manager, Proactive Care Lead, Southbourne Surgery |
| Shilan Ghafoor  | Medicines Governance Pharmacist, Medicines Governance, UKHSA |
| Michelle Jones | Principal Medicines Optimisation Pharmacist, Bristol North Somerset and South Gloucestershire Integrated Care Board |
| Elizabeth Luckett | Senior Screening and Immunisation Manager, Screening and Immunisation Team – Kent and Medway, NHSE South East |
| Dr Vanessa MacGregor | Consultant in Communicable Disease Control, East Midlands Health Protection Team, UKHSA  |
| Lesley McFarlane | Lead Immunisation Nurse Specialist, Immunisation Programmes Division, UKHSA |
| Briony Mason | Vaccination Manager and Professional Midwifery Advocate, Vaccination and Screening, NHSE West Midlands  |
| Tushar Shah | Lead Pharmacy Adviser, NHSE London  |

**2. Organisational authorisations**

This PGD is not legally valid until it has had the relevant organisational authorisation.

It is the responsibility of the organisation that has legal authority to authorise the PGD, to ensure that all legal and governance requirements are met. The authorising body accepts governance responsibility for the appropriate use of the PGD.

Insert authorising body name authorises this PGD for use by the services or providers listed below:

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| --- |
| Authorised for use by the following organisations and/or services |
| For instance, all NHS England commissioned immunisation services or NHS Trusts providing immunisation services.  |
| Limitations to authorisation |
| For instance, any local limitations the authorising organisation feels they need to apply in-line with the way services are commissioned locally. This organisation does not authorise the use of this PGD by ….  |

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| --- |
| Organisational approval (legal requirement) |
| Role | Name  | Sign | Date |
| For instance, NHS England Governance Lead, Medical Director |   |   |   |

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| Additional signatories according to locally agreed policy |
| Role | Name  | Sign | Date |
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Local enquiries regarding the use of this PGD may be directed to insert local contact.

[Section 7](#section7) provides a practitioner authorisation sheet. Individual practitioners must be authorised by name to work to this PGD. Alternative practitioner authorisation sheets may be used where appropriate in accordance with local policy but this should be an individual agreement or a multiple practitioner authorisation sheet as included at the end of this PGD.

#### Characteristics of staff

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| **Qualifications and professional registration**  | **All practitioners should only administer vaccinations where it is within their clinical scope of practice to do so. Practitioners must also fulfil the**[**additional requirements**](#AdditionalRequirements) **and** [**continued training requirements**](#Continued_training_req) **to ensure their competency is up to date, as outlined in the sections below**. Practitioners working to this PGD must also be one of the following registered professionals who can legally supply and administer under a PGD: * nurses and midwives currently registered with the Nursing and Midwifery Council (NMC)
* pharmacists and pharmacy technicians currently registered with the General Pharmaceutical Council (GPhC) (Note: This PGD is not relevant to privately provided community pharmacy services)
* dieticians, occupational therapists, paramedics, physiotherapists and podiatrists currently registered with the Health and Care Professions Council (HCPC)

Check [section 2](#section2) (Limitations to authorisation) to confirm whether all practitioners listed above have organisational authorisation to work under this PGD. |
| **Additional requirements** | Additionally, practitioners:* must be authorised by name as an approved practitioner under the current terms of this PGD before working to it
* must have undertaken appropriate training for working under PGDs for supply/administration of medicines
* must be competent in the use of PGDs (see [NICE Competency framework for healthcare professionals using PGDs](https://www.nice.org.uk/guidance/mpg2/resources))
* must be familiar with the vaccine product and alert to changes in the Summary of Product Characteristics (SPC), Immunisation Against Infectious Disease (the [Green Book](https://www.gov.uk/government/collections/immunisation-against-infectious-disease-the-green-book)), and national and local immunisation programmes
* must have undertaken training appropriate to this PGD as required by local policy and in line with the [[National Minimum Standards and Core Curriculum for Immunisation Training](https://www.gov.uk/government/publications/national-minimum-standards-and-core-curriculum-for-immunisation-training-for-registered-healthcare-practitioners)](https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/362171/National_Immun_Train_Stand1.pdf)
* must be competent to undertake immunisation and to discuss issues related to immunisation
* must be competent in the handling and storage of vaccines, and management of the cold chain
* must be competent in the appropriate administration method for the vaccine listed in this PGD
* must be competent in the recognition and management of anaphylaxis
* must have access to the PGD and associated online resources
* should fulfil any additional requirements defined by local policy

**Individual practitioners must be authorised by name, under the current version of this PGD before working according to it.** |
| **Continued training requirements**(continued over page) **Continued training requirements**(continued)  | Practitioners must ensure they are up to date with relevant issues and clinical skills relating to immunisation and management of anaphylaxis, with evidence of appropriate Continued Professional Development (CPD).Practitioners should be constantly alert to any subsequent recommendations from UKHSA, NHS England and other sources of medicines information. Note: The most current national recommendations should be followed, but a Patient Specific Direction (PSD) may be required to administer the vaccine in line with updated recommendations that are outside the criteria specified in this PGD. |

1. **Clinical condition or situation to which this PGD applies**

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| **Clinical condition or situation to which this PGD applies** | Shingrix® Herpes Zoster Vaccine (recombinant, adjuvanted) is indicated for the prevention of herpes zoster (‘zoster’ or shingles) and herpes zoster-related post-herpetic neuralgia (PHN) for adults who are eligible for the national shingles immunisation programme in accordance with the recommendations given in [Chapter 28a](https://www.gov.uk/government/publications/shingles-herpes-zoster-the-green-book-chapter-28a) of Immunisation Against Infectious Disease: the Green Book. |
| **Criteria for inclusion** | **Immunocompetent individuals** who: * reach the age of 65 **or** 70 years of age during the period 1 September 2023 and 31 August 2028. These individuals should be vaccinated on or after (but not before) their 65th or 70th birthday
* are aged between 70 years and 79 years old on or before 31 August 2023 **and** have never received a shingles vaccine
* are currently aged 80 years, but received a first dose of Shingrix® before turning 80 years old and require a second Shingrix® dose to complete the course before their 81st birthday

**Severely immunosuppressed[[2]](#footnote-3) individuals from 1 September 2025,** who**:** * are aged 18 years and above, and meet the definition of severe immunosuppression in the Box in [Chapter 28a](https://www.gov.uk/government/publications/shingles-herpes-zoster-the-green-book-chapter-28a). This includes individuals due to commence therapy. Note there is **no upper age limit**

**See** [**Appendix 1**](#Appendix2) **for further information on cohort eligibility**  |
| **Criteria for exclusion[[3]](#footnote-4)**(continued over page) **Criteria for exclusion[[4]](#footnote-5)**(continued)  | Individuals for whom no valid consent has been received (or for whom a best-interests decision in accordance with the [Mental Capacity Act 2005](https://www.legislation.gov.uk/ukpga/2005/9/contents), has not been obtained). For further information on consent, see [Chapter 2](https://www.gov.uk/government/publications/consent-the-green-book-chapter-2) of the Green Book. Several resources are available to inform consent (see [written information to be given to individual or carer](#writen_info) section). **Immunocompetent individuals on or after 1 September 2023** who:* have not yet reached their 65th birthday
* are between 65 to 69 years of age but reached their 65th birthday on or before 31 August 2023; these individuals must wait until they reach their 70th birthday
* are 80 years of age or over (except those who have received a partial course of Shingrix® who can receive a second dose to complete the course before their 81st birthday)

**All individuals** who: * have had a confirmed anaphylactic reaction to a previous dose of varicella-containing vaccine or to any other component of the shingles vaccine
* are suffering from acute severe febrile illness (the presence of a minor infection is not a contraindication for immunisation)
* have shingles infection with active lesions
* are pregnant
* have already received one dose of Zostavax® or 2 doses of Shingrix® prior to assessment. An exception is in place for individuals previously immunised with Zostavax® who subsequently become immunosuppressed (see [special considerations and additional information](#special_consi) section)
* have received a dose of Shingrix® in the last 8 weeks
* are aged under 18 years
 |
| **Cautions including any relevant action to be taken**  | Facilities for management of anaphylaxis should be available at all vaccination premises (see [Chapter 8](https://www.gov.uk/government/publications/vaccine-safety-and-adverse-events-following-immunisation-the-green-book-chapter-8) of the Green Book and advice issued by the [Resuscitation Council UK](https://www.resus.org.uk/about-us/news-and-events/rcuk-publishes-anaphylaxis-guidance-vaccination-settings)). The immunogenicity of the vaccine could be reduced in immunosuppressed subjects. However, vaccination should proceed in accordance with the national recommendations.Syncope (fainting) can occur following, or even before, any vaccination especially in adolescents as a psychogenic response to the needle injection. This can be accompanied by several neurological signs such as transient visual disturbance, paraesthesia and tonic-clonic limb movements during recovery. It is important that procedures are in place to avoid injury from faints. |
| **Action to be taken if the individual is excluded**(continued over page) **Action to be taken if the individual is excluded**(continued)  | Individuals who are not eligible for the national shingles immunisation programme should be advised when they will become eligible or why they are not eligible for immunisation. Refer to [Appendix 1](#Appendix1) for further information. If vaccination cannot be commenced before an immunocompetent individual is 80 years old, explain why vaccination will no longer be indicated.Individuals who have had a confirmed anaphylactic reaction to a previous dose of shingles vaccine or any components of the vaccine should be referred to a clinician for specialist advice and appropriate management. Individuals suffering from acute severe febrile illness should postpone immunisation until they have recovered. Advise when the individual may be vaccinated and ensure another appointment is arranged. When administration is postponed, arrange a future date for vaccination as appropriate, with due consideration of the individual’s age to ensure they will meet the inclusion criteria for immunisation.Individuals who present with a shingles infection with active lesions should postpone immunisation until recovered. As severely immunosuppressed individuals are at increased risk of recurrent zoster, Shingrix® can be given once any active shingles lesions have resolved.If clinically indicated, Shingrix® may be considered in pregnancy, following a discussion of the risks and benefits with the individual. In such cases, if the individual wishes to proceed with vaccination, Shingrix® may be given under a PSD. Immunocompetent individuals previously vaccinated with Zostavax® or who have received 2 doses of Shingrix® do not need to be revaccinated. Individuals who have been previously vaccinated with Zostavax® and who have since become severely immunosuppressed should be offered a course of Shingrix®, in line with other severely immunosuppressed individuals. See also [special considerations and additional information](#special_consi) section. A dose interval of 8 weeks is the shortest advised in [Chapter 28a](https://www.gov.uk/government/publications/shingles-herpes-zoster-the-green-book-chapter-28a), although this is [off label](#offlabel). The Shingrix® [SPC](http://www.medicines.org.uk) advises a dose interval of one to 2 months after the first dose for individuals who are presently or anticipated to become severely immunosuppressed. Administering the vaccine at an interval of less than 8 weeks is outside the scope of this PGD and a PSD should be used to administer the completing dose If required, seek advice from the local Screening and Immunisation Team, local Health Protection Team or the individual’s clinician as deemed appropriate. The risk to the individual of not being vaccinated must be taken into account.Document the reason for exclusion and any action taken in the individual’s clinical records.Inform or refer to the individual’s GP or a prescriber as appropriate. |
| **Action to be taken if the individual or carer declines treatment** | Informed consent, from the individual or a person legally able to act on the individual’s behalf, must be obtained prior to administration. Advise the individual or carer about the protective effects of the vaccine, the risks of infection and potential complications.Document advice given and the decision reached.Inform or refer to the individual’s GP or a prescriber as appropriate. |
| **Arrangements for referral for medical advice** | As per local policy |

1. **Description of treatment**

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| **Name, strength and formulation of drug** | Herpes zoster vaccine (recombinant, adjuvanted): * Shingrix®, powder and suspension for suspension for injection.

After reconstitution, one dose (0.5ml) of Shingrix® contains varicella zoster virus glycoprotein E antigen 50 micrograms, adjuvanted with AS01B. |
| **Legal category** | Prescription only medicine (POM). |
| **Black triangle** | No. |
| **Off-label use** | The Shingrix® [SPC](https://www.medicines.org.uk/emc/product/12054) advises an interval of 2 months between doses, which may be extended to between 2 and 6 months if flexibility is required. For individuals who are or about to become severely immunosuppressed and who might benefit from a shorter vaccination schedule, a one to 2 month interval between doses may be observed. The dose intervals advised in the Green Book and subsequently in this PGD are different to that outlined above; between **6 and 12 months for immunocompetent** individuals and 8 weeks to 6 months for severely immunosuppressed individuals. Vaccine should be stored according to the conditions detailed in the [storage](#Storage) section below. However, in the event of an inadvertent or unavoidable deviation of these conditions, refer to [Vaccine Incident Guidance](https://www.gov.uk/government/publications/vaccine-incident-guidance-responding-to-vaccine-errors). Where vaccine is assessed in accordance with these guidelines as appropriate for continued use this would constitute off-label administration under this PGD.Where a vaccine is recommended off-label, consider as part of the consent process, informing the individual or carer that the vaccine is being offered in accordance with national guidance but outside of product licence. |
| **Route and method of administration**(continued over page)**Route and method of administration**(continued) | Shingrix® must be reconstituted in accordance with the manufacturer’s instructions prior to administration. Following reconstitution, Shingrix® vaccine is given by intramuscular injection, preferably into the deltoid muscle of the upper arm.When administering at the same time as other vaccines, care should be taken to ensure that the appropriate route of injection is used for all the vaccinations. The vaccines should be given at separate sites, preferably into different limbs. If given into the same limb, they should be given at least 2.5cm apart. The site at which each vaccine was given should be noted in the individual’s records.Individuals with bleeding disorders may be vaccinated intramuscularly if, in the opinion of a clinician familiar with the individual's bleeding risk, vaccines or similar small volume intramuscular injections can be administered with reasonable safety by this route. Individuals on stable anticoagulation therapy, including individuals on warfarin who are up to date with their scheduled INR testing and whose latest INR was below the upper threshold of their therapeutic range, can be vaccinated via the intramuscular route. If the individual receives medication or other treatment to reduce bleeding, for example treatment for haemophilia, intramuscular vaccination can be scheduled shortly after such medication or treatment is administered. A fine needle (equal to 23 gauge or finer calibre such as 25 gauge) should be used for the vaccination, followed by firm pressure applied to the site (without rubbing) for at least 2 minutes. The individual or carer should be informed about the risk of haematoma from the injection. Subcutaneous administration is not recommended. Maladministration via the subcutaneous route may lead to an increase in transient local reactions. The vaccine should be visually inspected for foreign particulate matter and other variation of expected appearance before preparation and administration. Should either occur, do not administer the dose and discard the vaccine in accordance with local procedures. The reconstituted vaccine is an opalescent, colourless to pale brownish liquid. After reconstitution, the vaccine should be used promptly (see [storage](#Storage) section). The vaccine [SPC](https://www.medicines.org.uk/emc/product/12054) provides further guidance on preparation and administration. |
| **Dose and frequency of administration** | Single 0.5ml dose per administration. If the course is interrupted or delayed, it should be resumed as soon as possible, but not repeated. **Eligible immunocompetent individuals*** 2 doses of Shingrix®, with the second dose given **6 to 12 months** after the first dose.

**Severely immunosuppressed individuals (includes stem cell transplant recipients):** * 2 doses of Shingrix®, with the second dose given **8 weeks to 6 months** after the first dose

Severely immunosuppressed individuals who have already received 2 doses of Shingrix® do not require revaccination. See [special considerations and additional information](#special_consi) for managing individuals who subsequently become immunosuppressed after completing the shingles vaccine course.  |
| **Duration of treatment** | A 2 dose course (see [dose and frequency of administration](#DoseAndFrequencyOfAdministration)) |
| **Quantity to be supplied and administered** | Single 0.5ml dose per administration. |
| **Supplies** | Centrally purchased vaccines for the national immunisation programme can only be ordered via ImmForm and are provided free of charge.Protocols for the ordering, storage and handling of vaccines should be followed to prevent vaccine wastage (see Green Book [Chapter 3](https://www.gov.uk/government/publications/storage-distribution-and-disposal-of-vaccines-the-green-book-chapter-3)). |
| **Storage** | Store between +2°C to +8°C. Store in original packaging in order to protect from light. Do not freeze. Once reconstituted, any unused vaccine should be stored in a refrigerator at +2°C to +8°C if not immediately required. Though chemical and physical in-use stability has been demonstrated for 24 hours at +30°C, from a microbiological viewpoint, the reconstituted vaccine should be used as soon as possible. Any remaining vaccine should be discarded 6 hours following reconstitution.In the event of an inadvertent or unavoidable deviation of these conditions, vaccine that has been stored outside the conditions stated above should be quarantined and risk assessed on a case-by-case basis for suitability of continued off-label use or appropriate disposal. Refer to [Vaccine Incident Guidance](https://www.gov.uk/government/publications/vaccine-incident-guidance-responding-to-vaccine-errors).Contact the vaccine manufacturer where more specific advice is required about managing a temperature excursion.  |
| **Disposal**(continued over page) **Disposal**(continued)  | Follow local clinical waste policy and NHS standard operating procedures to ensure safe and secure waste disposal. Equipment used for immunisation, including used vials, ampoules, or discharged vaccines in a syringe or applicator, should be disposed of safely in a UN-approved puncture-resistant sharps box, according to local waste disposal arrangements and NHS England guidance [(HTM 07-01): safe and sustainable management of healthcare waste](https://www.england.nhs.uk/publication/management-and-disposal-of-healthcare-waste-htm-07-01/).  |
| **Drug interactions** | The immunological response may be diminished in individuals receiving immunosuppressant treatment. Vaccination is recommended even if the antibody response may be limited. See the [additional information](#Additionalinfo) section for information on co-administration with other vaccines.A detailed list of drug interactions is available in the [SPC](https://www.medicines.org.uk/emc/product/12054).  |
| **Identification and management of adverse reactions** | The most common adverse reactions observed after administration of Shingrix® are injection-site reactions (such as pain, redness and swelling) myalgia and headache. Most of these reactions are not long-lasting (median duration of 2 to 3 days). Other very common side-effects include gastrointestinal symptoms (including nausea, vomiting, diarrhoea and abdominal pain), chills and fever.In the event of a person developing a varicella (widespread) or shingles-like (dermatomal) rash post-Shingrix® vaccination, a vesicle fluid sample should be sent for analysis, to confirm the diagnosis and determine whether the rash is vaccine-associated or wild-type (see [Chapter 28a](https://www.gov.uk/government/publications/shingles-herpes-zoster-the-green-book-chapter-28a)). A detailed list of adverse reactions is available in the [SPC](https://www.medicines.org.uk/emc/product/12054). |
| **Reporting procedure of adverse reactions** | Healthcare professionals, individuals and carers are encouraged to report suspected adverse reactions to the Medicines and Healthcare products Regulatory Agency (MHRA) using the [Yellow Card reporting scheme](http://yellowcard.mhra.gov.uk/) or by searching for MHRA Yellow Card in the Google Play or Apple App Store.Any adverse reaction to the vaccine should be documented in the individual’s record and the individual’s GP should be informed.  |
| **Written information to be given to patient or carer** | Offer the marketing authorisation holder’s patient information leaflet (PIL) provided with the vaccine.Immunisation promotional material may be provided as appropriate. [Shingles vaccination guide](https://www.gov.uk/government/publications/shingles-vaccination-for-adults-aged-70-or-79-years-of-age-a5-leaflet)[Shingles easy-read leaflet](https://www.healthpublications.gov.uk/ViewProduct.html?sp=Sshingleseasyreadleaflet)For resources in accessible formats and alternative languages, please visit [Home- Health Publications](https://www.healthpublications.gov.uk/). Where applicable, inform the individual or carer that large print, Braille or audio CD PILs may be available from emc accessibility (freephone 0800 198 5000) by providing the medicine name and product code number, as listed on the [SPC](https://www.medicines.org.uk/emc/xpil#gref). |
| **Advice and follow-up treatment** | Inform the individual or carer of possible side effects and their management.Give advice regarding normal reactions to the injection, for example redness and pain at the injection site.The individual or carer should be advised to seek medical advice in the event of a severe adverse reaction and report this via the [Yellow Card Scheme](http://yellowcard.mhra.gov.uk/). When administration is postponed, advise the individual or carer when to return for vaccination with due consideration of the individual’s age to ensure they will meet the inclusion criteria for immunisation. If vaccination cannot be commenced before an immunocompetent individual is 80 years old, explain why vaccination will no longer be indicated (there is no upper age limit for severely immunocompromised individuals). Individuals should be advised to seek medical attention if they develop a varicella (widespread) or shingles-like (dermatomal) rash post-Shingrix® vaccination. |
| **Special considerations and additional information** | Ensure there is immediate access to adrenaline (epinephrine) 1 in 1000 injection and easy access to a telephone at the time of vaccination.**Dosing intervals**[Chapter 28a](https://www.gov.uk/government/publications/shingles-herpes-zoster-the-green-book-chapter-28a) advises a minimum interval of 8 weeks between doses for individuals. For operational reasons, **a recommended interval of 6 months between doses is in place for immunocompetent individuals**. **This PGD cannot be used to administer Shingrix® to immunocompetent individuals at an interval of less than 6 months**.Individuals anticipated to commence immunosuppressive treatment should ideally be assessed before starting treatment. The first dose of Shingrix® should be given at least 2 weeks before treatment starts, though one month prior is preferred. If immunosuppressive treatment is subsequently commenced after the first dose of Shingrix® is given, the second dose may be given 8 weeks to 6 months later**.** **Administration with other vaccines**Shingrix® can be given at the same time as unadjuvanted inactivated influenza vaccines or 23-valent pneumococcal vaccine (PPV23). The vaccines should be administered at different injection sites. The adverse reactions of fever and shivering are more frequent when PPV23 vaccine is co-administered with Shingrix®.Whilst clinical data on co-administration of the 20-valent pneumococcal conjugate vaccine (PCV20) and Shingrix® is limited, both vaccines may be given together. The individual should be made aware that side effects may be more potentiated. In line with general advice about co-administration of inactivated vaccines, Shingrix® can be given concomitantly with inactivated influenza vaccines. Initially, a 7 day interval was recommended between Shingrix® and adjuvanted influenza vaccine because the potential reactogenicity from 2 adjuvanted vaccines may reduce tolerability in those being vaccinated. Interim data from a US study on co-administration of Shingrix® with adjuvanted seasonal influenza vaccine is reassuring. Therefore, an appointment for administration of the seasonal influenza vaccine can be an opportunity to also provide shingles vaccine, although the latter should be offered all year round, rather than purely as a seasonal programme. A 7 day gap between administration of Shingrix® and COVID-19 vaccine is no longer required. As Shingrix® is an inactivated vaccine, where individuals in an eligible cohort present having received another inactivated or live vaccine, Shingrix® vaccination should still be considered. In most cases, vaccination should proceed to avoid any further delay in protection and to avoid the risk of the individual not returning for a later appointment. In such circumstances, individuals should be informed about the likely timing of potential adverse events relating to each vaccine.**Individuals who subsequently become immunosuppressed after completing a shingles vaccine course** Most individuals who have been previously vaccinated with either one dose of Zostavax® or 2 doses of Shingrix® do not require a repeat course of Shingrix®. A clinical exception is in place for individuals who become severely immunosuppressed subsequent to receiving a dose of Zostavax®. In line with the inclusion criteria, a 2 dose course of Shingrix® should be offered to these individuals. The need for booster doses of Shingrix® has not yet been determined, so a repeat course of Shingrix® should not be offered where such individuals have already completed a Shingrix® course (see [Chapter 28a](https://www.gov.uk/government/publications/shingles-herpes-zoster-the-green-book-chapter-28a)).  |
| **Records**  | The practitioner must ensure the following is recorded: * that valid informed consent was given, or a decision to vaccinate was made in the individual’s best interests in accordance with the [Mental Capacity Act 2005](https://www.legislation.gov.uk/ukpga/2005/9/contents)
* name of individual, address, date of birth and GP with whom the individual is registered
* name of immuniser
* name and brand of vaccine
* date of administration
* dose, form and route of administration of vaccine
* quantity administered
* batch number and expiry date
* anatomical site of vaccination
* advice given, including advice given if the individual is excluded or declines immunisation
* details of any adverse drug reactions and actions taken
* supplied via PGD

Records should be signed and dated (or password-controlled on e-records). All records should be clear, legible and contemporaneous.This information should be recorded in the individual’s GP record and any other appropriate medical records, such as care or nursing records. When the vaccine has been administered to individuals under 19 years of age, notify the local Child Health Information Service (CHIS) using the appropriate documentation or pathway as required by any local or contractual agreement. A record of all individuals receiving treatment under this PGD should also be kept for audit purposes in accordance with local policy. |

1. **Key references**

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| **Key references**  | **Shingles*** Shingrix® Summary of Product Characteristics. GlaxoSmithKline UK, updated 3 March 2025 <https://www.medicines.org.uk/emc/product/12054/smpc>
* Immunisation Against Infectious Disease: The Green Book, [Chapter 28a](https://www.gov.uk/government/publications/shingles-herpes-zoster-the-green-book-chapter-28a), updated July 2023

<https://www.gov.uk/government/collections/immunisation-against-infectious-disease-the-green-book> * DHSC: JCVI statement on the shingles (herpes zoster) vaccination programme, published 13 November 2024 <https://www.gov.uk/government/publications/shingles-herpes-zoster-vaccination-programme-jcvi-statement-november-2024/jcvi-statement-on-the-shingles-herpes-zoster-vaccination-programme#expansion-of-eligibility-immunosuppressed-people>
* Shingles: Guidance and Vaccination Programme. Updated 4 July 2023.

<https://www.gov.uk/government/collections/shingles-vaccination-programme> * UKHSA: Vaccination against shingles- information for healthcare practitioners, updated 6 July 2023 <https://www.gov.uk/government/publications/shingles-vaccination-guidance-for-healthcare-professionals>

**General*** NHSE Health Technical Memorandum 07-01: safe and sustainable management of healthcare waste, updated 7 March 2023 <https://www.england.nhs.uk/publication/management-and-disposal-of-healthcare-waste-htm-07-01>
* National Minimum Standards and Core Curriculum for Immunisation Training, published 7 February 2018 <https://www.gov.uk/government/publications/national-minimum-standards-and-core-curriculum-for-immunisation-training-for-registered-healthcare-practitioners>
* NICE Medicines Practice Guideline 2 (MPG2): Patient Group Directions, last updated 27 March 2017 <https://www.nice.org.uk/guidance/mpg2>
* NICE MPG2 Patient group directions: competency framework for health professionals using patient group directions, updated 4 January 2018

<https://www.nice.org.uk/guidance/mpg2/resources> * UKHSA Immunisation Collection. <https://www.gov.uk/government/collections/immunisation>
* Vaccine Incident Guidance: responding to errors in vaccination storage, handling and administration, updated 7 July 2022

<https://www.gov.uk/government/publications/vaccine-incident-guidance-responding-to-vaccine-errors>  |

1. **Practitioner authorisation sheet**

**Shingles (Shingrix®) PGD v3.0 Valid from: 1 September 2025 Expiry: 31 August 2028**

Before signing this PGD, check that the document has had the necessary authorisations in section 2. Without these, this PGD is not lawfully valid.

**Practitioner**

By signing this PGD, you are indicating that you agree to its contents and that you will work within it.

PGDs do not remove inherent professional obligations or accountability.

It is the responsibility of each professional to practise only within the bounds of their own competence and professional code of conduct.

|  |
| --- |
| I confirm that I have read and understood the content of this PGD and that I am willing and competent to work to it within my professional code of conduct. |
| Name | Designation | Signature | Date |
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**Authorising manager**

|  |
| --- |
| I confirm that the practitioners named above have declared themselves suitably trained and competent to work under this PGD. I give authorisation on behalf of insert name of organisation for the above named health care professionals who have signed the PGD to work under it. |
| Name | Designation | Signature | Date |
|  |  |  |  |

**Note to authorising manager**

Score through unused rows in the list of practitioners to prevent practitioner additions post managerial authorisation.

This authorisation sheet should be retained to serve as a record of those practitioners authorised to work under this PGD.

**Appendix 1: Immunocompetent patients and phased implementation of Shingrix® during Stage 1 (1 September 2023 to 31 August 2028)**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Implementation stage** | **Age attained (years) within first programme year (1 Sept 2023 to 31 Aug 2024)** | **Dates of birth** | **Programme year offered** | **Age offered at** |
| Previously eligible cohorts: | 70 to 79 | before 1 Sept 1953 | those in the cohort previously eligible for the Zostavax® programme prior to 1 Sept 2023 remain eligible for Shingrix® up to their 80th birthday. Completing doses must be given before the individual’s 81st birthday |
| Individuals reaching the age of 70 between 1 September 2023 and 31 August 2028  | 70 | 1 Sept 1953 to 31 Aug 1954 | Year 1 | 1 Sept 2023 to 31 Aug 2024 | 70 |
| 69 | 1 Sept 1954 to 31 Aug 1955 | Year 2 | 1 Sept 2024 to 31 Aug 2025 | 70 |
| 68 | 1 Sept 1955 to 31 Aug 1956 | Year 3 | 1 Sept 2025 to 31 Aug 2026 | 70 |
| 67 | 1 Sept 1956 to 31 Aug 1957 | Year 4 | 1 Sept 2026 to 31 Aug 2027 | 70 |
| 66 | 1 Sept 1957 to 31 Aug 1958 | Year 5 | 1 Sept 2027 to 31 Aug 2028 | 70 |
| Individuals reaching the age of 65 between 1 September 2023 and 31 August 2028  | 65 | 1 Sept 1958 to 31 Aug 1959 | Year 1 | 1 Sept 2023 to 31 Aug 2024 | 65 |
| 64 | 1 Sept 1959 to 31 Aug 1960 | Year 2 | 1 Sept 2024 to 31 Aug 2025 | 65 |
| 63 | 1 Sept 1960 to 31 Aug 1961 | Year 3 | 1 Sept 2025 to 31 Aug 2026 | 65 |
| 62 | 1 Sept 1961 to 31 Aug 1962 | Year 4 | 1 Sept 2026 to 31 Aug 2027 | 65 |
| 61 | 1 Sept 1962 to 31 Aug 1963 | Year 5 | 1 Sept 2027 to 31 Aug 2028 | 65 |

1. This includes any relevant amendments to legislation [↑](#footnote-ref-2)
2. Immunocompromised individuals in this PGD, are those defined as severely immunosuppressed, as outlined in the Box in [Chapter 28a](https://www.gov.uk/government/publications/shingles-herpes-zoster-the-green-book-chapter-28a) [↑](#footnote-ref-3)
3. Exclusion under this PGD does not necessarily mean the medication is contraindicated, but it would be outside its remit and another form of authorisation will be required [↑](#footnote-ref-4)
4. Exclusion under this PGD does not necessarily mean the medication is contraindicated, but it would be outside its remit and another form of authorisation will be required [↑](#footnote-ref-5)